US ERA ARCHIVE DOCUMENT

### DATA EVALUATION RECORD ALGAE OR DIATOM EC50 TEST GUIDELINE 122-2 OR 123-2 (TIER I OR II)

1. CHEMICAL: 2-chloro-4,6-bis(isopropylamino)-s-triazine

<u>Code No.</u>: 080808

TEST MATERIAL: Propazine Purity: 98%

3. CITATION

Authors: D. W. Gledhill; J. B. Bussard

Title: Acute toxicity of propazine to Anabaena

flos-aquae

Study Completion Date: 5/22/95

Laboratory: ABC Laboratories, Inc.
Sponsor: Griffin Corporation
Laboratory Report ID: ABC Labs #41968-9

DP Barcode: D237791 MRID No.: 442873-12

REVIEWED BY: Thomas M. Steeger, Ph.D., Fishery Biologist, EEB, ERB IV, U.S. EPA

Signature: Thomas M. Streger

Date: /0/2/97

Nicholas E. Federoff, Wildlife Biolgist, EFED, APPROVED BY:

ERB IV, U.S. EPA

Signature:

STUDY PARAMETERS

Scientific Name of Test Organism: Anabaema flos-aquae

Definitive Test Duration: 120 hours

Type of Concentrations: Mean measured/Nominal

CONCLUSIONS: This study is scientifically sound and does fulfill the 123-2 guideline requirements for acute toxicity tests for algae. The 120-hour EC50 was estimated to be 0.18 mg a.i./L. After 120 hours, the no-observed effect concentration was 0.068 mg a.i./L.

Results Synopsis

EC<sub>50</sub>: 0.18 ppm ai NOEL: '0.068 ppm ai 95% C.I.: 0.16-0.20 ppm ai

Slope: 3.64

8. ADEQUACY OF THE STUDY

Classification: Core

B. Rationale: Methodology was consistent with FIFRA guidelines. Conclusions were verifiable.



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- C. Repairability:

### 9. GUIDELINE DEVIATIONS

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## 10. SUBMISSION PURPOSE:

#### 11. MATERIALS AND METHODS

# A. Test Organisms

Guideline Criteria	Reported Information
Species Skeletonema costatum Anabaena flos-aquae Selenastrum capricornutum Navicula pelliculosa	Anabaena flos-aquae
Initial Number of Cells 3,000 - 10,000 cells/ml	$4.1 \times 10^3 \text{ cells/ml}$
Nutrients Standard formula, e.g. 20XAAP	macro/micronutrient stock solutions

## B. Test System

Guideline Criteria	Reported Information	
Solvent	dimethylformamide (DMF)	
Temperature Skeletonema: 20°C Others: 24-25°C	24 ± 2°C	
Light Intensity  Anabaena: 2.2 K lux (±15%) Others: 4.3 K lux (±15%)	4,310 ± 650 Lux	
Photoperiod Skeletonema: 14 h light, 10 h dark or 16 h light, 8 h dark Others: Continuous	Continuous	

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	Guideline Criteria Reported Information
Hq	Range: 7.5 - 7.8
	keletonema: approx. 8.0 thers: approx. 7.5

# C. Test Design

Guideline Criteria	Reported Information	
Dose range 2X or 3X progression	2X	
<u>Doses</u> at least 5	control, solvent control, 0.065, 0.13, 0.25, 0.50, and 1.0 mg/L	
Controls negative and/or solvent	Control and solvent control	
Replicates per dose 3 or more (4 or more for Navicula)	triplicates	
Duration of test 120 hours	120 hours	
Daily observations were made?	Yes	
Method of Observations	hemacytometer/Olympus Model BH-2 microscope	
Maximum Labeled Rate	1.2 lb ai/acre	

# 12. REPORTED RESULTS

Guideline Criteria	Reported Information
Quality assurance and GLP compliance statements were included in the report?	Yes
Initial and 120 h cell densities were measured?	Yes (in control and solvent control)
Control cell count at 120 hr ≥2X initial count?	Yes (512X)

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Guideline Criteria	Reported Information
Initial chemical concentrations measured? (Optional)	Yes
Raw data included?	Yes

Dose Response

Dose (mg ai/L)	Cell Density (x 104 cells/ml)	% Inhibition	120-Hour pH
Control	210		7.8
Solvent Control	210		7.8
0.068	220	+4.76	7.8
0,13	160	-23.81	7.7
0.26	42	-80.91	7.7
0.52	32	-85.45	77
1.0	8.9	-95.95	7.6

### Other Significant Results:

### Statistical Results

Statistical Method: ANOVA (Proc GLM)/multiple means comparison (Dunnett's); Proc NLIN used to estimate B and EC50

EC<sub>50</sub>: 0.18 ppm

95% C.I.: 0.16 - 0.20 ppm

Slope: 3,64

NOEC: 0.068 ppm

## 13. Verification of Statistical Results

Statistical Method: TOXANAL

EC<sub>50</sub>: 0.201 ppm

95% C.I.: 0.101-0.370 ppm

Slope: 3.12

NOEC: 0.068 ppm

Adjusted for active ingredient:

EC<sub>50</sub>: 0.18 ppm ai 95% C.I.: 0.16 - 0.20 ppm ai

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NOEC: 0.068 ppm ai

14. REVIEWER'S COMMENTS: